



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C11D 17/00, 3/37, 17/06, 3/18, 3/12, 3/22, 11/00, 10/04, 1/94 // 1:04, 1:12, 1:90	A1	(11) International Publication Number: WO 99/24546 (43) International Publication Date: 20 May 1999 (20.05.99)
(21) International Application Number: PCT/EP98/06989 (22) International Filing Date: 22 October 1998 (22.10.98) (30) Priority Data: 08/964,617 5 November 1997 (05.11.97) US (71) Applicant (for AU BB CA CY GB GD GH GM IE IL KE LC LK LS MN MW NZ SD SG SL SZ TT UG ZW only): UNILEVER PLC [GB/GB]; Unilever House, Blackfriars, London EC4P 4BQ (GB). (71) Applicant (for all designated States except AU BB CA CY GB GD GH GM IE IL KE LC LK LS MN MW NZ SD SG SL SZ TT UG ZW): UNILEVER N.V. [NL/NL]; Weena 455, NL-3013 AL Rotterdam (NL). (72) Inventors: FARRELL, Terence, James; Unilever Research U.S. Inc., 45 River Road, Edgewater, NJ 07020 (US). SHAFER, Georgia, Lynn; Unilever Research U.S. Inc., 45 River Road, Edgewater, NJ 07020 (US). DALTON, James, Joseph; 7009 E. Acoma Drive #2041, Scottsdale, AZ 85254 (US). HE, Mengtao; Unilever Research U.S. Inc., 45 River Road, Edgewater, NJ 07020 (US). MCFANN, Gregory, Jay; Unilever Research U.S. Inc., 45 River Road, Edgewater, NJ 07020 (US).	(74) Agent: MOLE, Peter, Geoffrey; Unilever plc, Patent Division, Colworth House, Sharnbrook, Bedford MK44 1LQ (GB). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	

(54) Title: BAR COMPOSITION COMPRISING ENTRAPPED EMOLLIENT DROPLETS DISPERSED THEREIN

(57) Abstract

The present invention relates to novel chip compositions comprising specifically defined carriers; benefit agent; and thickening agent (e.g., fumed silica). In a second embodiment, the invention comprises bar compositions comprising mixtures of the chips of the invention and chips containing defined surfactant systems. The invention further comprises a process for forming the chips of the invention and a method of enhancing deposition of benefit agent without compromising processing using the chips of the invention.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Larvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

- 1 -

BAR COMPOSITION COMPRISING ENTRAPPED
EMOLLIENT DROPLETS DISPERSED THEREIN

5 The present invention relates to bar compositions,
particularly synthetic soap bar compositions, able to
deliver beneficial agents (e.g., silicone, petrolatum,
maleated soybean oil) in higher amounts than previously
possible. In particular, the invention relates to bar
10 compositions comprising emollient droplets entrapped in a
thickened carrier. The emollient containing thickened
carrier compositions are formed as separate chip/powder
compositions and are then mixed with "base" chips
(comprising the surfactant system) prior to milling,
15 extruding and stamping the bars. The invention further
relates to a method of making the additives. The invention
further relates to a method of enhancing delivery of benefit
agent from bars which method comprises mixing the base chips
with the benefit agent containing chips, milling, extruding
20 and stamping.

It has long been a desirable goal to deliver some kind
of benefit agent (e.g., silicone or other oils) to the skin
through a personal wash composition.

25

In liquid cleansers, for example, cationic hydrophilic
polymers such as Polymer JR^(R) from Amerchol or Jaguar^(R)
from Rhone Poulenc have been used to enhance delivery of
benefit agents (EP 93,602; WO 94/03152; and WO 94/03151).
30 In the applicants' copending application, U.S. Serial No.
08/412,803 to Tsaur et al., separate hydrogel particles act

- 2 -

as a structure to entrap the benefit agent in concentrated form.

In the present invention, entrapment of benefit agent is achieved by producing compositions comprising emollient wherein the medium in which the emollients are found (e.g., hydrophobically modified polyalkylene glycols; polyoxyethylene - polyoxypropylene copolymers; or mixture of either or both with polyalkylene glycols) is thickened with a thickening agent (e.g., fumed silica) so that the droplets are entrapped in the medium. More specifically, and without wishing to be bound by theory, the above-identified compounds are believed to act as a matrix entrapping the emollient, and the thickener is believed to stop emollient from escaping from the matrix. In addition, mixing hydrophobically modified polyalkylene glycol (PAGs) or EO-PO copolymers with polyalkylene glycol can fine-tune the dissolution rate of adjuvant compounds used to make the bar so that they dissolve at same rate as surfactant chips also used in making the bar.

The emollient-containing thickened carrier compositions may be inserted into bars in concentrated forms (as in the applicants' copending application No. 08/828,443 to Rattinger et al.) or dispersed throughout the bar. The thickened carrier composition should be prepared separate from the base bar composition and the separate chip compositions should be coextruded.

Delivery of benefit agents (e.g., silicone, maleated soybean oil) in bar compositions has proven more difficult in bars for a number of reasons. If the benefit agent is added homogeneously (i.e., is intimately mixed with other

- 3 -

components), for example, the generally hydrophobic benefit agent is in intimate contact with hydrophobic materials in the base composition and is not "available" for deposition. Thus, little or no benefit agent will be present in the
5 final bar (after milling, plodding and extrusion of chips) to be delivered to the skin. If the benefit agent is too viscous, on the other hand, it tends to get in the processing equipment and become too difficult to process.

10 U.S. Patent No. 5,154,849 to Visscher et al. teaches bar compositions containing a silicone skin mildness/moisturizing aid component. In one embodiment, the silicone component may be mixed with a carrier which is selected to facilitate incorporation of the silicone.

15 Preferred carrier is said to be polyethylene glycol. At column 16, the reference describes that silicone is mixed into melted Carbowax (polyethylene glycol), that the mixture is cooled to form flakes, and that the flakes are preferably added to an amalgamator.

20 It is clear, however, that the Visscher et al. reference contemplates a silicone/carrier system different from the benefit agent/carrier/thickener system of the subject invention. First, the Visscher patent does not
25 teach a thickener (e.g., fumed silica or water soluble starch), a critical component of the emollient containing compositions and one which is believed to provide the structure required to retain and engulf the benefit agent (e.g., silicone or other benefit agents) in the carrier.

30 Secondly, as suggested above and as shown in Figures 1 and 2, the structure of the carrier/silicone chip is distinct. The Visscher et al. composition does not contain the silicone in discrete droplets, but rather the silicone oozes

- 4 -

and surrounds the carrier. By contrast, the benefit agent droplets of the invention are discrete droplets retained within the chip. This helps to ensure the silicone benefit agent does not ooze and interfere with processing.

5

The discrete particles of the invention, in turn, are present for two reasons, it is believed. The first, as noted above, is presence of thickening agent (e.g. water soluble starch or fumed silica) which, while not wishing to be bound by theory, it is believed helps to thicken the carrier (e.g., hydrophobically modified polyalkylene glycol; EO-PO copolymers; mixtures of one or both with polyalkylene glycol) such that the viscosity of the carrier is minimum 800 centipoise (cps), preferably greater than 1500 cps, more preferably greater than 3000 cps and can thereby entrap the benefit agent. The second reason is that, unlike the Visscher et al. system, the present invention requires there be an equal amount or more of carrier relative to the benefit agent. By contrast, it appears from Visscher et al., where eleven pounds of silicone (column 15, lines 1-2), are mixed with 5 to 6 pounds of Carbowax (column 15, line 29) that there is probably an excess of silicone to PEG and, at the least, there is no recognition of the criticality of having an equal amount or more of carrier to benefit agent.

25

In short, the chips of the Visscher reference are extremely difficult to process both because there is no control over the amount of silicone used, and also because there is no use of thickened carrier.

30

Finally, Visscher teaches polyalkylene glycol carrier, but does not teach or suggest use of hydrophobically

- 5 -

modified polyalkylene glycols; EO-PO copolymers; or mixtures of one or both with polyalkylene glycols.

In one embodiment of the present invention, the applicants have unexpectedly found that, when specific additive compositions are made containing an equal amount or greater of carrier (e.g., hydrophobically modified polyalkylene glycol; EO-PO copolymers; mixtures of one or both these groups with polyalkylene glycols) to benefit agent, and further containing a thickening agent for the carrier such that the viscosity of carrier is 800 cps or greater, preferably greater than 1500 cps, more preferably greater than 3000 cps, the benefit agent (e.g., silicone, petrolatum, maleated soybean oil etc.) becomes entrapped as discrete droplets in the thickened carrier, which in turn allows the benefit agent to be much more readily processed.

Specifically, in this embodiment the invention comprises a chip composition comprising:

- (a) 40% to about 80% by wt. of the chip composition of a carrier selected from:
- (1) hydrophobically modified polyalkylene glycol having MW of about 4,000-25,000 wherein the compound has formula $(AG)_m-R$ or $R-(AG)_m-R$, AG being alkylene glycol monomer unit, m being greater than 50 and R being attached hydrophobic group;
 - (2) a polyoxyethylene-polyoxypropylene nonionic copolymer having MW about 4000 to 25,000;
 - (3) mixtures of (1) and (2); and
 - (4) mixtures of (1) and/or (2) with polyalkylene glycol having a molecular weight greater than

- 6 -

- 4000, preferably greater than 5,000 to
20,000, more preferably 5000 to 10,000;
- (b) 10% to 40% by wt. of the chip composition of
benefit agent (e.g., silicone petrolatum, maleated
5 soybean oil);
- (c) 0.01% to 30% by wt. chip composition thickening
agent;
- (d) 0% to 10% by wt. chip composition, preferably 0%
to 5% by wt. water; and
- 10 (e) 0% to 15% by wt. chip composition
structurant/filler selected from C₈ to C₂₄ fatty
acid or ester, C₈ to C₂₄ alcohol or ether
derivative, or mixtures thereof. Preferably, it
is a C₈ to C₂₄ straight chain, saturated fatty
15 acid.

The invention further comprises an extruded bar
composition which is produced using about 5 to 50%,
preferably 10 to 40%, more preferably 20 to 40% chips as
20 described above and about 95% to 50% chips comprising about
5% to 90% by wt. of a surfactant system wherein the
surfactant is selected from soap, anionic surfactant,
nonionic surfactant, amphoteric surfactant, zwitterionic
surfactant, cationic surfactant and mixtures thereof. The
25 "soap and/or surfactant" chips additionally may comprise
other components typically found in such chips such as, for
example, minor amounts of fragrance, preservative (e.g.,
butylated hydroxy toluene) skin feel polymer (e.g., guar)
etc. It may also contain free fatty acid and/or
30 structurant/inert filler.

- 7 -

Although the surfactant system of the second chip may be a pure soap surfactant system, preferably the surfactant system comprises:

- (a) a first synthetic surfactant which is an anionic surfactant; and
- (b) a second synthetic surfactant selected from the group consisting of a second anionic different from the first, a nonionic, an amphoteric and mixtures thereof.

A particularly preferred surfactant system comprises acyl isethionate as the first anionic and a sulfosuccinate or a betaine surfactant or mixtures of the two.

In a further embodiment of the invention, the invention comprises a method of making benefit agent containing chips comprising:

- (a) 40% to 80% of a carrier selected from one of groups (a) (1) - (4) above;
- (b) 10% to 40% benefit agent;
- (c) 0.01% to 30% thickening agent;
- (d) 0% to 10% water; and
- (e) 0% to 10% structurant/filler which can be a C₈ to C₂₄ fatty acid or ester derivative or C₈ to C₂₄ alcohol or ether derivative, wherein the method comprises mixing the ingredients at temperatures above the melting point of the carrier (i.e., above about 50°C) for 1 to 60 minutes; cooling on a chill roll (at about 0° to 25°C); and collecting.

- 8 -

The invention will now be further described by way of example only with reference to the accompanying figures, in which:

5 Figure 1 is a micrograph of a chip produced following process described in U.S. patent No. 5,154,849 to Visscher et al. wherein the chip is flooded with water and photographed under a microscope. The silicone does not form discrete particles but forms a large layer of silicone
10 surrounding the polyalkylene glycol. The viscous silicone gets caught in machinery and inhibits processing; and

 Figure 2 is a micrograph of a chip product according to the present invention (i.e., forming benefit agent chip,
15 forming surfactant chip, and mixing together). As noted, the benefit agent is found in discrete drops. Further, it is obvious that there is far less benefit agent present (e.g., to interfere with processing) than in Figure 1.

20 In one embodiment of the invention, the present invention relates to novel soap chip compositions (e.g., in the process for making bars, molten compositions are formed which are then cooled on what is commonly called a chill roll to form flakes or chips; these chips are subsequently
25 refined and/or plodded to form billets which are stamped and cut to form final bars) which are readily processable in conventional soap machinery, while still showing significant benefit agent deposition (i.e., comparable to deposition obtained in liquid body washes).

30

 As seen from Figures 1 and 2, by carefully controlling the level of benefit agent (so that it cannot exceed the level of carrier) and by utilizing thickening agent, such

- 9 -

as, for example, starches or fumed silica (while not wishing to be bound by theory, it is believed the thickening agent thickens the carrier such that the emollient is entrapped in the carrier), the applicants have been able to provide
5 discrete droplets of benefit agent so that the agent is unable to stick to the machinery and significantly inhibit processing. Further, the emollient/benefit agent more readily deposits from the bar.

10 The emollient is prepared in one chip/composition and, base bar is separately prepared, and chips are later mixed. This is described below.

With regard to the separate chip composition, the first
15 component (carrier) of the emollient chip may be a hydrophobically modified polyalkylene glycol (HMPAG) having broad molecular weight 4,000 to 25,000, preferably 4,000 to 15,000.

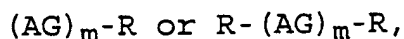
20 Generally, the polymers will be selected from polyalkylene glycols chemically and terminally attached by hydrophobic moieties, wherein the hydrophobic moiety can be derivatives of linear or branched alkyl, aryl, alkylaryl, alkylene, acyl (e.g., having a carbon number of C₂ to C₆₀,
25 preferably C₈ to C₄₀; fat and oil derivatives of alkylglyceryl, glyceryl, sorbitol, lanolin oil, coconut oil, jojoba oil, castor oil, almond oil, peanut oil, wheat germ oil, rice bran oil, linseed oil, apricot pits oil, walnuts, palm nuts, pistachio nuts, sesame seeds, rapeseed, cade oil,
30 corn oil, peach pit oil, poppyseed oil, pine oil, soybean oil, avocado oil, sunflower seed oil, hazelnut oil, olive oil, grapeseed oil, and safflower oil, Shea butter, babassu

- 10 -

oil, etc. These hydrophobically modified polyalkylene glycols are usually commercially available (see Table 1 for examples).

5 To ensure water solubility, it is preferred that the portion of alkylene oxide moiety per mole of HMPAG is between 60% wt. and 99% wt. (preferably 85% wt. to 97% wt.). In other words, the total content of the hydrophobic moiety is between 1% wt. and 40% wt. (preferably 3% wt. to 15% wt.)
10 per mole of the defined HMPAG.

In general, the HMPAGs of the invention have the following formula:



15

where AG is the alkylene glycol monomer unit (generally ethylene or propylene glycol), and m is greater than 50. R is any of the hydrophobic moieties described above.

20 Specifically, examples of various hydrophobically modified polyalkylene glycols are set forth in Table 1 below where in T_m ($^{\circ}\text{C}$) were obtained from literature from the corresponding chemical suppliers or measured by the inventors using a differential scanning calorimetry
25 technique.

Table 1: Representative hydrophobically modified PEGs.

(R = Hydrophobic moieties such as linear or branched alkyl chains (e.g., having carbon number of C4 to C40);

30 derivatives of sorbitol, lanolin radical, coconut radical, jojoba acid radical, castor oil radical, etc.;

POE = Polyoxyethylene (e.g., $-(\text{CH}_2\text{CH}_2\text{O})_m\text{H}$);

- 11 -

m = No. ethylene oxide monomer units; m>50).

Chemicals	Suppliers (Brands)	Comments
POE(m)-R solid.	Witco (Varonic LI-420)	R = glyceryltallowate; m = 200; white
solid.	Seppic (Simusol 220Tm)	R = glycerylstearate; m = 200; white
solid.	Americhol (glucam E-200)	R = glucoside; m = 200; white water soluble; white solid
	Calgene Chemical (600-S)	Tm:52-62C; R = stearate; m = 150; Tm:52-62C
	Calgene Chemical (600-L)	R = laurate; m = 150
R-POE (m)-R	Stepan (KESSCO PEG6000 distearate)	R = stearate; m = 174; Tm:54C; white solid

As noted, melting temperature of the compounds is preferred to be about 25°-85°.

The carrier may also be a polyoxyethylene polyoxypropylene nonionic copolymer (EO-PO) copolymers.

The polyoxyethylene polyoxypropylene nonionic copolymers (EO-PO copolymers) of the subject invention are generally commercially available polymers having a broad molecular weight range and EO/PO ratio and a melting temperature of from about 25° to 85°C, preferably 40° to 65°C.

Generally, the polymers will be selected from one of two classes of polymers, i.e., (1) (EO)_m(PO)_n(EO)_m type copolymers or (PO)_n(EO)_m(PO)_n type copolymers of defined m/n ratio and optional hydrophobic moieties (e.g., decyltetradecanol ether) attached to either EO or PO compounds (such products are commercially available for example, from BASF under the Trademark Pluronic^(R) or

- 12 -

Pluronic-R^(R), respectively); or (2) EO-PO polymers with amine constituents such as $N_2C_2H_4(PO)_{4n}(EO)_{4m}$ or $N_2C_2H_4(EO)_{4m}(PO)_{4n}$ with defined values of m and n and optional hydrophobic moieties attached to either EO or PO components (such products are commercially available, for example from BASF as Tetronic^(R) and Tetronic-R^(R), respectively).

Specifically, examples of various Pluronic and Tetronic EO-PO polymers are set forth in Table 1 below wherein T_m (°C) and Ross Miles foam height data (measured at 0.1% and 50°C) were digested from literature from BASF.

- 13 -

TABLE 1

Polymer		$T_m(^{\circ}\text{C})$	Foam Heights (ml)	EO and PO Number
Pluronic:	$(\text{EO})_m - (\text{PO})_n - (\text{EO})_m$			m/n
	F38	48	35	46/16
	F68	52	35	75/30
	F77	48	47	52/35
	F87	49	44	62/39
	F88	54	48	97/39
	F98	58	43	122/47
	F108	57	41	128/54
	F127	56	41	98/67
Pluronic-R:	$(\text{PO})_n - (\text{EO})_m - (\text{PO})_n$			
	10R8	46	20	90/9
	17R8	53	2	155/15
	25R8	54	15	227/21
Tetronic:	$\text{N}_2\text{C}_2\text{H}_4 - (\text{PO})_{4n}(\text{EO})_{4m}$			
	707	46	60	35/12
	1107	51	50	64/20
	908	58	40	85/16
	1307	54	40	78/25
	1508	60	40	159/30
Tetronic-R:	$\text{N}_2\text{C}_2\text{H}_4 - (\text{EO})_{4m}(\text{PO})_{4n}$			
	90R8	47	0	90/17
	110R7	47	0	64/21
	150R8	53	0	12/29

- 14 -

In general, the molecular weight of the copolymers used ranges from 2,000 to 25,000 (preferably 3,000 to 10,000).

5 The EO-terminated polymers (Pluronic and Tetronic) are preferred to the PO-terminated ones (Pluronic-R and Tetronic-R) for the advantages of mildness enhancement and lather generation. To ensure water solubility, it is preferred that the portion of ethylene oxide moiety per mole
10 is between 50% to 90% wt., more preferably 60-85% wt. In other words, 2m:n (for Pluronic) or m:n (for Tetronic) ranges from 1.32 to 11.9, preferably 2.0 to 7.5.

As noted, melting temperature of the compounds must be
15 about 25°-85°, preferably 40° to 65°C, the latter being more favorable for processing (e.g., chips form more easily and logs plod more readily).

Finally, the carrier can be mixtures of hydrophobically
20 modified PAG with EO-PO copolymers; mixtures of hydrophobically modified PAG with polyalkylene glycols; mixtures of EO-PO copolymers with polyalkylene glycol or mixtures of both HMPAG and EO-PO copolymers with polyalkylene glycol, wherein polyalkylene glycol is defined
25 as having a MW greater than 4000 to about 100,000, preferably 4000 to 10,000. An especially preferred carrier is polyethylene glycol, for example, Carbowax PEG 8000^(R) from Union Carbide.

30 One advantage of using mixtures of either or both of HMPAG and EO-PO copolymers with polyalkylene glycol is to fine-tune dissolution rate of said chip composition to be same as that of surfactant chips. This is important for bar

- 15 -

user properties (e.g., in maintaining bar integrity during wash).

Finally, it should be noted that use of polyalkylene glycol alone is also contemplated as carrier for this invention (with any benefit agent and any thickener) and that this is disclosed in applicant's copending U.S. Serial No. 08/828,442.

10 The benefit agent of the subject invention may be a single benefit agent component or it may be a benefit agent compound added via a carrier. Further the benefit agent composition may be a mixture of two or more compounds one or all of which may have a beneficial aspect. In addition, the benefit agent itself may act as a carrier for other components one may wish to add to the bar composition.

The benefit agent can be an "emollient oil" by which is meant a substance which softens the skin (stratum corneum) by increasing into water content and keeping it soft by retarding decrease of water content.

Preferred emollients include:

- 25 (a) silicone oils, gums and modifications thereof such as linear and cyclic polydimethylsiloxanes; amino, alkyl alkylaryl and aryl silicone oils;
- (b) fats and oils including natural fats and oils such as jojoba, soybean (including maleated soybean oil), rice bran, avocado, almond, olive, sesame, persic, castor, coconut, mink oils; cacao fat; beef tallow, lard; hardened oils obtained by hydrogenating the aforementioned oils; and
- 30 synthetic mono, di and triglycerides such as

- 16 -

myristic acid glyceride and 2-ethylhexanoic acid glyceride;

(c) waxes such as carnauba, spermaceti, beeswax, lanolin and derivatives thereof;

5 (d) hydrophobic plant extracts;

(e) hydrocarbons such as liquid paraffins, vaseline, microcrystalline wax, ceresin, squalene, pristan and mineral oil;

10 (f) higher fatty acids such as lauric, myristic, palmitic, stearic, behenic, oleic, linoleic, linolenic, lanolic, isostearic and poly unsaturated fatty acids (PUFA);

(g) higher alcohols such as lauryl, cetyl, stearyl, oleyl, behenyl, cholesterol and 2-hexydecanol alcohol;

15 (h) esters such as cetyl octanoate, myristyl lactate, cetyl lactate, isopropyl myristate, myristyl myristate, isopropyl palmitate, isopropyl adipate, butyl stearate, decyl oleate, cholesterol isostearate, glycerol monostearate, glycerol

20 distearate, glycerol tristearate, alkyl lactate, alkyl citrate and alkyl tartrate;

(i) essential oils such as mentha, jasmine, camphor, white cedar, bitter orange peel, ryu, turpentine,

25 cinnamon, bergamot, citrus unshiu, calamus, pine, lavender, bay, clove, hiba, eucalyptus, lemon, starflower, thyme, peppermint, rose, sage, menthol, cineole, eugenol, citral, citronelle, borneol, linalool, geraniol, evening primrose,

30 camphor, thymol, spirantol, penene, limonene and terpenoid oils;

- 17 -

- (j) lipids such as cholesterol, ceramides, sucrose esters and pseudo-ceramides as described in European Patent Specification No. 556,957;
- (k) vitamins such as vitamin A and E, and vitamin alkyl esters, including those vitamin C alkyl esters;
- (l) sunscreens such as octyl methoxyl cinnamate (Parsol MCX) and butyl methoxy benzoylmethane (Parsol 1789);
- (m) phospholipids; and
- (n) mixtures of any of the foregoing components.

A particularly preferred benefit agent is silicone, preferably silicones having viscosity greater than about 10,000 centipoise. The silicone may be a gum and/or it may be a mixture of silicones. One example is polydimethylsiloxane having viscosity of about 60,000 centistokes. Other preferred emollients include petrolatum, maleated soybean oil and sunflower seed oil.

The benefit agent generally comprises about 10% to 40%, preferably 20% to 40%, most preferably 25% to 40% by weight of the chip composition.

A criticality of the invention is the presence of a thickening agent which is believed required to thicken the viscosity of the carrier.

The thickening agent must thicken the carrier such that the thickened carrier has a viscosity of at least 800 centipoises (cps), preferably at least 1500 cps, most preferably greater than 3000 cps.

- 18 -

Examples of thickening agents which may be used include silicas and starches. Among the starches which may be used are water soluble starches such as maltodextrin or partially soluble starches such as potato or corn starch. By water
5 soluble is meant that a 10% by wt. or greater solution of the starch in water will dissolve to form a clear or substantially clear solution (except for small amounts of insoluble residue which may impart a translucent haziness to otherwise clear solution).

10

A particularly prepared thickening agent is fumed silica. Fumed silica is generally produced by the hydrolysis of silicon tetrachloride vapor in a flame of hydrogen and oxygen. The process produces particles of from
15 about 7 to 30 millimicrons.

The enormous surface area and chain forming abilities are believed to allow it to form three-dimensional networks, altering flowing properties i.e., cause thickening.

20

The thickening agent will generally comprise the 0.01 to 30% by wt. of the composition, preferably 5% to 20% by wt., most preferably 5% to 10% by wt. of the composition.

25

It should be noted when fumed silica is used, thickener should comprise no more than about 10%.

In terms of optional components, water comprises 0 to 10%, preferably 0% to 8% by wt., most preferably 0.1 to 5%
30 by wt. of the chip composition. It is sometimes preferred to have little or no additional water (other than that inherently present in the compounds) in the chip mixture because this may sometimes cause processing difficulties.

- 19 -

In addition the chip composition may comprise 0% to 15%, preferably 2% to 10% fatty acid, i.e., C₈ to C₂₄ fatty acid. Generally, this is a straight chain, saturated fatty acid although this is not necessarily the case. The fatty acid helps to modify the wear rate of the emollient chip to better match that of the base soap.

The chip may also comprise a structuring aid and/or filler which can be fatty acid as described above or ester derivative; or a preferably straight and saturated C₈ to C₂₄ alcohol or ether derivative.

In terms of the base bar compositions, the invention comprises extruded bar compositions in which 5% to about 50%, preferably 10% to 40%, more preferably 20% to 40% of the chips used to make the final bars comprise the benefit agent additives (i.e. chips) described above and in which 95% to 50%, preferably 90% to 60%, most preferably 80% to 60% of the chips comprise chips which comprise the surfactant system defining the final bar.

Specifically, the surfactant system chips comprise about 5% to 90% by wt. of a surfactant system wherein the surfactant is selected from soap (pure soap surfactant systems are included), anionic surfactant, nonionic surfactant, amphoteric zwitterionic surfactant, cationic surfactant and mixtures thereof. These chips may additionally comprise other components typically found in final bar compositions, for example, minor amounts of fragrance, preservative, skin feel polymer etc.

- 20 -

In terms of the surfactant system, the term "soap" is used herein in its popular sense, i.e., the alkali metal or alkanol ammonium salts of aliphatic alkane- or alkene monocarboxylic acids. Sodium, potassium, mono-, di- and tri-ethanol ammonium cations, or combinations thereof, are suitable for purposes of this invention. In general, sodium soaps are used in the compositions of this invention, but from about 1% to about 25% of the soap may be potassium soaps. The soaps useful herein are the well known alkali metal salts of natural or synthetic aliphatic (alkanoic or alkenoic) acids having about 12 to 22 carbon atoms, preferably about 12 to about 18 carbon atoms. They may be described as alkali metal carboxylates of acrylic hydrocarbons having about 12 to about 22 carbon atoms.

Soaps having the fatty acid distribution of coconut oil may provide the lower end of the broad molecular weight range. Those soaps having the fatty acid distribution of peanut or rapeseed oil, or their hydrogenated derivatives, may provide the upper end of the broad molecular weight range.

It is preferred to use soaps having the fatty acid distribution of coconut oil or tallow, or mixtures thereof, since these are among the more readily available fats. The proportion of fatty acids having at least 12 carbon atoms in coconut oil soap is about 85%. This proportion will be greater when mixtures of coconut oil and fats such as tallow, palm oil, or non-tropical nut oils or fats are used, wherein the principle chain lengths are C_{16} and higher. Preferred soap for use in the compositions of this invention has at least about 85% fatty acids having about 12 to 18 carbon atoms.

- 21 -

Coconut oil employed for the soap may be substituted in whole or in part by other "high-alluric" oils, that is, oils or fats wherein at least 50% of the total fatty acids are composed of lauric or myristic acids and mixtures thereof. These oils are generally exemplified by the tropical nut oils of the coconut oil class. For instance, they include: palm kernel oil, babassu oil, ouricuri oil, tucum oil, cohune nut oil, murumuru oil, jaboty kernel oil, khakan kernel oil, dika nut oil, and ucuhuba butter.

A preferred soap is a mixture of about 15% to about 20% coconut oil and about 80% to about 85% tallow. These mixtures contain about 95% fatty acids having about 12 to about 18 carbon atoms. The soap may be prepared from coconut oil, in which case the fatty acid content is about 85% of C₁₂-C₁₈ chain length.

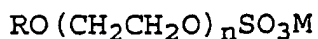
The soaps may contain unsaturation in accordance with commercially acceptable standards. Excessive unsaturation is normally avoided.

Soaps may be made by the classic kettle boiling process or modern continuous soap manufacturing processes wherein natural fats and oils such as tallow or coconut oil or their equivalents are saponified with an alkali metal hydroxide using procedures well known to those skilled in the art. Alternatively, the soaps may be made by neutralizing fatty acids, such as lauric (C₁₂), myristic (C₁₄), palmitic (C₁₆), or stearic (C₁₈) acids with an alkali metal hydroxide or carbonate.

- 22 -

The anionic detergent active which may be used may be aliphatic sulfonates, such as a primary alkane (e.g., C₈-C₂₂) sulfonate, primary alkane (e.g., C₈-C₂₂) disulfonate, C₈-C₂₂ alkene sulfonate, C₈-C₂₂ hydroxyalkane sulfonate or alkyl glyceryl ether sulfonate (AGS); or aromatic sulfonates such as alkyl benzene sulfonate.

The anionic may also be an alkyl sulfate (e.g., C₁₂-C₁₈ alkyl sulfate) or alkyl ether sulfate (including alkyl glyceryl ether sulfates). Among the alkyl ether sulfates are those having the formula:

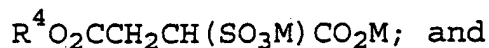


wherein R is an alkyl or alkenyl having 8 to 18 carbons, preferably 12 to 18 carbons, n has an average value of greater than 1.0, preferably greater than 3; and M is a solubilizing cation such as sodium, potassium, ammonium or substituted ammonium. Ammonium and sodium lauryl ether sulfates are preferred.

The anionic may also be alkyl sulfosuccinates (including mono- and dialkyl, e.g., C₆-C₂₂ sulfosuccinates); alkyl and acyl taurates, alkyl and acyl sarcosinates, sulfoacetates, C₈-C₂₂ alkyl phosphates and phosphates, alkyl phosphate esters and alkoxyl alkyl phosphate esters, acyl lactates, C₈-C₂₂ monoalkyl succinates and maleates, sulphoacetates, alkyl glucosides and acyl isethionates.

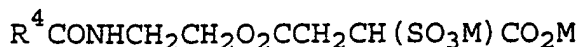
- 23 -

Sulfosuccinates may be monoalkyl sulfosuccinates having the formula:



5

amide-MEA sulfosuccinates of the formula;

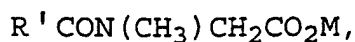


10

wherein R^4 ranges from C_8 - C_{22} alkyl and M is a solubilizing cation.

Sarcosinates are generally indicated by the formula:

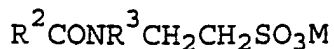
15



wherein R^1 ranges from C_8 - C_{20} alkyl and M is a solubilizing cation.

20

Taurates are generally identified by formula:



wherein R^2 ranges from C_8 - C_{20} alkyl, R^3 ranges from C_1 -
25 C_4 alkyl and M is a solubilizing cation.

Particularly preferred are the C_8 - C_{18} acyl isethionates.

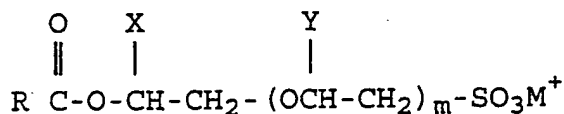
These esters are prepared by reaction between alkali metal isethionate with mixed aliphatic fatty acids having from 6
30 to 18 carbon atoms and an iodine value of less than 20. At

- 24 -

least 75% of the mixed fatty acids have from 12 to 18 carbon atoms and up to 25% have from 6 to 10 carbon atoms.

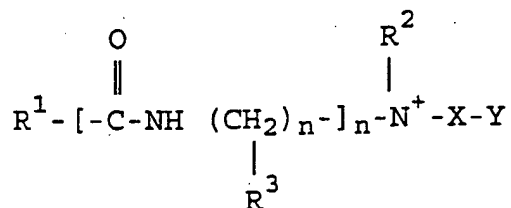
Acyl isethionates, when present, will generally range from about 10% to about 70% by weight of the total bar composition. Preferably, this component is present from about 30% to about 60%.

The acyl isethionate may be an alkoxylated isethionate such as is described in Ilardi et al., U.S. Patent No. 5,393,466, hereby incorporated by reference. This compound has the general formula:



wherein R is an alkyl group having 8 to 18 carbons, m is an integer from 1 to 4, X and Y are hydrogen or an alkyl group having 1 to 4 carbons and M⁺ is a monovalent cation such as, for example, sodium, potassium or ammonium.

Amphoteric detergents which may be used in this invention include at least one acid group. This may be a carboxylic or a sulphonic acid group. They include quaternary nitrogen and therefore are quaternary amido acids. They should generally include an alkyl or alkenyl group of 7 to 18 carbon atoms. They will usually comply with an overall structural formula:



- 25 -

where R^1 is alkyl or alkenyl of 7 to 18 carbon atoms;

R^2 and R^3 are each independently alkyl, hydroxyalkyl or
5 carboxyalkyl of 1 to 3 carbon atoms;

m is 2 to 4;

n is 0 to 1;

10

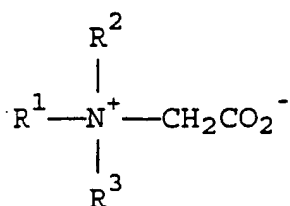
X is alkylene of 1 to 3 carbon atoms optionally
substituted with hydroxyl, and

Y is $-\text{CO}_2-$ or $-\text{SO}_3-$

15

Suitable amphoteric detergents within the above general
formula include simple betaines of formula:

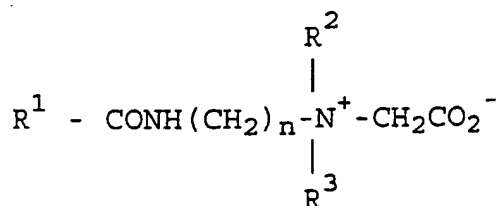
20



25

and amido betaines of formula:

30



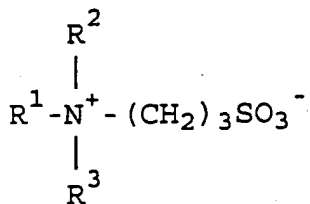
35

where n is 2 or 3.

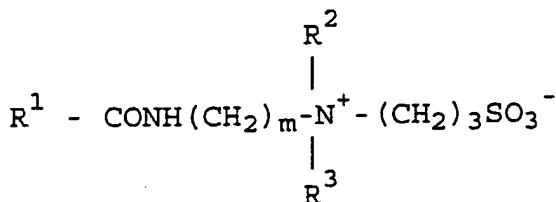
- 26 -

In both formulae R^1 , R^2 and R^3 are as defined previously. R^1 may in particular be a mixture of C_{12} and C_{14} alkyl groups derived from coconut so that at least half, preferably at least three quarters of the groups R^1 have 10
5 to 14 carbon atoms. R^2 and R^3 are preferably methyl.

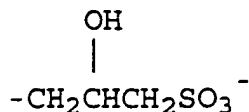
A further possibility is that the amphoteric detergent is a sulphobetaine of formula:



or



where m is 2 or 3, or variants of these in which $-(CH_2)_3 SO_3^-$ is replaced by



In these formulae R^1 , R^2 and R^3 are as discussed previously.

The nonionic which may be used as the second component of the invention include in particular the reaction products of compounds having a hydrophobic group and a reactive
35 hydrogen atom, for example aliphatic alcohols, acids, amides

- 27 -

or alkylphenols with alkylene oxides, especially ethylene oxide either alone or with propylene oxide. Specific nonionic detergent compounds are alkyl (C₆-C₂₂) phenols ethylene oxide condensates, the condensation products of
5 aliphatic (C₈-C₁₈) primary or secondary linear or branched alcohols with ethylene oxide, and products made by condensation of ethylene oxide with the reaction products of propylene oxide and ethylenediamine. Other so-called
10 nonionic detergent compounds include long chain tertiary amine oxides, long chain tertiary phosphine oxides and dialkyl sulphoxides.

The nonionic may also be a sugar amide, such as a polysaccharide amide. Specifically, the surfactant may be
15 one of the lactobionamides described in U.S. Patent No. 5,389,279 to Au et al. which is hereby incorporated by reference or it may be one of the sugar amides described in Patent No. 5,009,814 to Kelkenberg, hereby incorporated into the subject application by reference.

20

Examples of cationic detergents are the quaternary ammonium compounds such as alkyldimethylammonium halogenides.

25

Other surfactants which may be used are described in U.S. Patent No. 3,723,325 to Parran Jr. and "Surface Active Agents and Detergents" (Vol. I & II) by Schwartz, Perry & Berch, both of which are also incorporated into the subject application by reference.

30

- 28 -

Although the bar may be a pure soap bar, preferably the surfactant system of this chip (forming the surfactant system in the bar) comprises:

- (a) a first synthetic surfactant which is anionic; and
- 5 (b) a second synthetic surfactant selected from the group consisting of a second anionic different from the first, a nonionic, an amphoteric and mixtures thereof.

10 The first anionic can be any of those recited above, but is preferably a C₈ to C₁₈ isethionate as discussed above. Preferably acyl isethionate will comprise 10% to 90% by wt. total bar composition.

15 The second surfactant is preferably a sulfosuccinate, a betaine or mixtures of the two. The second surfactant or mixture of surfactant will generally comprise 1% to 10% total bar composition. A particularly preferred composition comprises enough sulfosuccinate to form 3-8% total bar
20 compositions and enough betaine to form 1-5% of total bar composition.

The base bar composition may also comprise water and structurant/filler as described in connection with the chip
25 composition (e.g., fatty acids or esters, alcohols or ethers thereof). The structurant may also be polyalkylene glycol with molecular weight between 2,000 and 20,000, preferably 3000 and 10,000. Such PEGs are commercially available, such as those marketed under tradename PEG 8000^(R) or PEG 4000^(R)
30 from Union Carbide.

- 29 -

Other ingredients that can be used as structurants or fillers include starches, preferably water soluble starches such as maltodextrin and polyethylene wax or paraffin wax.

5 Structuring aids can also be selected from water soluble polymers chemically modified with hydrophobic moiety or moieties, for example, EO-PO block copolymer, hydrophobically modified PEGs such as POE(200-glyceryl-stearate, glucam DOE 120 (PEG Methyl Glucose Dioleate), and
10 Hodg CSA-102 (PEG-150 stearate), and Rewoderm^(R) (PEG modified glyceryl cocoate, palmate or tallowate) from Rewo Chemicals.

Other structuring aids which may be used include
15 Amerchol Polymer HM 1500 (Nonoxynyl Hydroethyl Cellulose).

Finally, bars of the invention may comprise 0% to 25%, preferably 2% to 15% by wt. of an emollient such as ethylene glycol, propylene glycol and/or glycerine. Small amounts of
20 these emollients can be added to base bar to modify lather attributes, skin feel etc.

In relation to processing, in general, the additive, benefit agent chips are formed by mixing the ingredients in
25 a mixer at a temperature just above the melting point of the polyalkylene glycol (e.g., about 50°C and above, generally no higher than about 110°C) for about 1 to 60 minutes, and then cooling in a chill roll. Order of addition is not critical. The "non" benefit agent chips are formed by similarly mixing
30 and cooling (If used in one mixer, the same ranges and temperatures are used).

- 30 -

The chips are then combined, for example, in a hopper or ribbon mixer where they may be refined (e.g., worked into a more pliable mass), plodded into billets, stamped and cut.

5 In a further embodiment of the invention, the invention relates to a method of forming additives (chips) containing a benefit agent which method comprises:

- 10 (a) mixing carrier, benefit agent, thickener, optional water and optional fatty acid in a container for 1 to 60 minutes at about above 50°C; and
- (b) cooling the mixture on a chill roll to about 0 to 25°C to form chips.

The following examples are intended to further
15 illustrate the invention and are not intended to limit the invention in any way.

Unless stated otherwise, all percentages are intended to be percentages by weight.

20

EXAMPLES

Protocol

25 Silicone measurement was conducted as follows:

Analysis is done by method known as ICP (Inductively Coupled Argon Plasma). This procedure required a step involving extraction with xylene, and is therefore currently
30 used only in-vitro. The ICP technique employed a Thermo Jarrell Ash Atom Scan 25 with measurements being made at 251.612 nm. Additional ICP measurement parameters are given below.

- 31 -

The treatment process was as follows:

Porcine skin was shaved, dermatomed, and sectioned into 25 cm pieces prior to treatment. The skin sample was then
5 treated by rubbing the bar sample across the skin 10 times, in a back and forth motion. The resulting liquor on the skin was lathered for 30 seconds and then rinsed for 10 seconds with water which was regulated at 32-35°C (90-95F). The treated skin sample was placed in a borosilicate
10 scintillation vial that contained 10 ml of xylene. The samples were placed on a platform shaker for 1 hour to allow for the extraction of the silicone. After the extraction period, the skin was removed from the vial and the extract was analyzed using ICP technique. Sample solutions were
15 tested against a 10 pm silicone standard.

What is measured is deposition of silicone (or other emollient) in parts per million.

20 Typical ICP Measurement Parameters for Measuring
 Silicone in Xylene

Torch gas flow	high
Auxiliary gas flow	1.5 L/min
Analyzer pump rate	0.9 m L/min
Nebulizer pressure	21 psi
Observation height	12 mm above load cell
Plasma power	1750 W
Wavelength	251.612 nm
Slit height	6 mm
Integration time	4 sec

- 32 -

Example 1

Using the protocol discussed above, benefit agent
5 deposition (e.g., deposition of silicone) was measured in
compositions representing (1) the bar of Visscher et al.
with no fumed silica chips; (2) the bars of the invention
which did contain fumed silica chips; and (3) a liquid body
wash composition. Each is discussed in greater detail
10 below:

(1) Visscher Bar (WO 92/08444)

The Visscher bar was obtained following the procedure
15 taken from WO 92/08444 (equivalent to U.S. Patent No.
5,154,849) where polyethylene glycol is used as a carrier
for silicone in bars (procedure was done in a Patterson
mixture). The procedure was as follows:

20 (a) 681 gm of Carbowax PEG 8000 was melted and held
around 60°C;

(b) 400 gm of GE 350 cps silicone was added; and

(c) 273 gm of GE 500,000 cps silicone was added.

25 (The patent explains the carrier to be 10:9 silicone
A:PEG where silicone A is a blend of 40:60 silicone gum,
500,000 cps to silicone fluid, 350 cps)

The mixture remained in the mixer for 45 minutes until
30 it was considered homogenous. The mixture was then removed
and placed on a chill roll set at 7°C. The resulting "chips"
were soft, pliable and severely tacky. Silicone covered the
entire surface of the equipment.

- 33 -

A sample bar was prepared by chip mixing surfactant chip: Visscher chip ratio of 4:1 (wherein surfactant chip comprises 40-60% fatty acid isethionate, 20-30% fatty acid, 5 1-10% sodium isethionate, 1-10% sulfosuccinate, about 5% betaine, preservatives, dyes and minors); and extruding into a billet with a Weber Selander plodder. The resulting billet was soft and from experience not considered a viable product. The pressed bar lathered poorly. From experience 10 this type of "chip" cannot be produced using conventional equipment.

More specifically, mixing surfactant chips and Visscher chips at a weight ratio of 4:1, respectively, resulted in 15 large, non-free flowing clumps which adhered together by surface silicone. This result impeded feeding into the extruder. Material which did feed was extruded as a soft, sticky billet. When stamped, the bar had a poor surface, was tacky and produced little lather when wetted.

20

(2) Bar of the Invention

The bar of the invention comprised a 70%/30% mixture of chips wherein the 30% additive chip component had the 25 following formulation range:

40-100%, preferably 40-80% polyethyleneglycol (e.g. PEG 8000);
10-50%, preferably 10-40% polydimethyl siloxane of 60,000 centistokes;
30 0.1 to 10%, preferably 1 to 5% Cab-o-sil^(R) fumed silica (e.g., fumed silica 45-5);
0-20%, preferably 1-10% deionized water; and

- 34 -

0-20%, preferably 0-10% to C₈ to C₂₂ fatty acid

and

the 70% surfactant chips were like the surfactant chips used in the Visscher et al. bar, as follows:

- 5 about 40-60% by wt. fatty acid isethionate;
about 20-30% by wt. fatty acid;
about 1-10% by wt. sodium isethionate
about 1-10% by wt. sulfosuccinate;
about 5% by wt. betaine; and
10 remainder preservative, dyes, water and other minors.

A preferred benefit agent chip comprises as follows:

- (a) 55-65% PEG
(b) 25-40% silicone
15 (c) 1-7% fumed silica; and
(d) 0-8% deionized water.

The chips were mixed, plodded together at the above-identified ratios, and extruded into bars.

20

(3) Liquid Body Wash

The liquid body wash had the following formulation:

25

	% by wt.
Betaine	5-15%
Sodium Cocoyl Isethionate	1-10%
Anionic	1-5%
Fragrance, preservatives	0.1-2.0%
Water	to balance

- 35 -

As noted deposition results were taken using the ICP techniques discussed and results set forth as follows:

	Deposition.
Visscher Bar	2.16 +/- 0.48 $\mu\text{g}/\text{cm}^2$
Bar of Invention	2.24 +/- 0.83 $\mu\text{g}/\text{cm}^2$
Liquid	2.14 +/- 0.62 $\mu\text{g}/\text{cm}^2$

5 It is surprising that the bar can deposit as well as the liquids. Moreover, in contrast to Visscher, the bar of the invention was readily processable and did not clog machinery (See Example 2).

10 Example 2

To further show differences between the bar of the invention and bars of Visscher, applicants decided to analyze the chips more closely.

15 Chips used in formation of the Visscher et al. bar, and chips carrying benefit agent and used in the formation of the bars of the invention were micrographed.

20 As seen from Figures 1 and 2, the Visscher et al. (P&G) chips show large "blobs" of silicone surrounding the alkylene glycol while the chips of the invention show small discrete droplets of silicone.

25 While not wishing to be bound by theory, it is believed the difference in amount of silicone and how it is formed accounts for the tremendous processing difficulties experienced in forming the P&G bars relative to those of the

- 36 -

invention. As noted above, 4:1 ratio of Visscher chips to surfactant chips formed large non-free flowing clumps which hindered chip feeding into the extruder and noodle processing. The clumps also caused agglomeration in the
5 vacuum chamber which significantly reduced billet formation. Further, as noted, material which did extrude was soft and sticky and, when stamped, the bar had a poor surface, was tacky and produced little lather when wetted.

- 37 -

Example 3

A chip composition having the following composition was prepared:

5

852 g PEG 8000 (polyethylene glycol w/ MW of about 8000)	46.3%
426 g EO-PO copolymer*	23.2%
526 g maleated soybean oil	28.6%
35 g fumed silica	1.9%

* Pluronic F108: (EO)₁₂₈ (PO)₅₄ (EO)₁₂₈

The composition was prepared as follows:

10

PEG 8000 and EO-PO were melted up in overhead mixer and were allowed to deaerate. Maleated soybean oil and fumed silica were stirred in. After 2 minutes dispersion, the mixture was poured onto chill roll and collected as solid flakes. Melt temperature was 85°C (185F).

15

This example shows both mixtures of EO-PO copolymer and polyalkylene (helping fine-tune dissolution of chips to similar of that of surfactant chips); and also shows use of maleated soybean oil.

20

Example 4

A chip composition having the following composition was prepared:

25

- 38 -

1000 g PEG 8000	51.3%
500 g EO-PO copolymer*	25.6%
320 g petrolatum	16.4%
91 g sunflower seed oil	4.7%
38 g fumed silica	2.0%

* Pluronic F108: (EO)₁₂₈ (PO)₅₄ (EO)₁₂₈

5 The composition was prepared as follows:

Petrolatum was premixed with sunflower oil to make it liquid. PEG 8000 and EO-PO were melted up in overhead mixer and allowed to deaerate. Petrolatum/sunflower oil mixture
10 was stirred in, followed by fumed silica. The mixture was poured onto chill roll. Melt temperature was 84°C(183F).

Example 5

15 A chip composition having the following composition was prepared:

1000 g PEG 8000	49.0%
hydrophobically modified PEG*	24.5%
250 g petrolatum	12.2%
250 g PDMS, 100,000 cSt	12.2%
40 g fumed silica	2.0%

20 * POE (200) glyceryl stearate

The composition was prepared as follows:

- 39 -

PEG 8000 and hydrophobically modified were melted in overhead mixer. Petrolatum, PDMS, and fumed silica were added. The mixture was poured onto chill roll.

5

This example shows mixture of hydrophobically modified polyalkylene glycol and alkylene glycol as well as petrolatumas benefit agent.

10 Example 6

A chip composition having the following composition was prepared:

1000 g PEG 8000	45.3%
500 g EO-PO copolymer*	22.7%
662 g PDMS, 100,000 cSt	30.0%
44 g fumed silica	1.8%

15

*Pluronic F108: (EO)₁₂₈ (PO)₅₄ (EO)₁₂₈

Chips were prepared as in Examples 3-5.

20

Example 7

1000 g of each of the chips of Examples 3-6 (representing 34% w/w of final bar) were combined with 1941 g of Dove^(R) as surfactant chips (representing 66% w/w of final bar) in a ribbon blender and extruded into bars in a standard manner. The Dove^(R) surfactant chips had composition as follows:

25

- 40 -

about 40-60% by wt. fatty acid isethionate;
about 20-30% by wt. fatty acid;
about 1-10% by wt. sodium isethionate.
5 about 1-10% by wt. sulfosuccinate;
about 5% by wt. betaine; and
remainder preservative, dyes, water and other minors

Throughput rate was as good as using chips of Dove^(R)
10 alone. Further, the rheological properties were comparable
to that of Dove^(R). These experiments showed that the
emollient containing chips could be successfully incorporated
into bars without affecting processing and thus the emollient
can be subsequently successfully delivered. As noted, a
15 broad range of emollient oils can be delivered.

- 41 -

CLAIMS

1. A benefit chip composition comprising:

(a) 40% to about 80% by wt. of chip composition of a carrier selected from the group consisting of:

(1) hydrophobically modified polyalkylene glycol having MW of about 4000 - 25,000, wherein the compound has formula

(AG)_m-R or R-(AG)_m-R,

wherein AG is alkylene glycol monomer unit, m > 50 and R is attached hydrophilic group;

(2) polyoxyethylene-polyoxypropylene copolymer having MW about 4000 to 25,000;

(3) mixtures of (1) and (2); and

(4) mixture of (1) and/or (2) with polyalkylene glycol having MW greater than 4000 to 20,000.

(b) 10% to 40% by wt. chip composition benefit agent;

(c) 0.01 to 10% by wt. chip composition thickener;

(d) 0 to 10% by wt. chip composition water; and

(e) 0% to 15% by wt. chip composition, i.e.,

structuring aid/filler selected from C₈ to C₂₄

fatty acids or ester derivatives, and C₈ to C₂₄

alcohols or ether derivatives, and mixtures thereof,

wherein the viscosity of the thickened carrier is equal

to or greater than 800 cps.

2. A composition according to claim 1, wherein the MW of alkylene carrier used in mixture (3) is 5,000 to 10,000.

3. A composition according to claim 1 or claim 2, wherein benefit agent comprises silicone.

- 42 -

4. A composition according to any of the preceding claims, wherein benefit agent comprises petrolatum.

5. A composition according to any of the preceding claims, wherein benefit agent comprises maleated soybean oil.

6. A composition according to any of the preceding claims, wherein benefit agent comprises sunflower seed oil.

7. A composition according to any of the preceding claims, wherein the thickener comprises fumed silica.

8. A composition according to any of the preceding claims, wherein thickener comprises a water soluble starch.

9. A composition according to any of the preceding claims, wherein viscosity is at least 1500 cps.

10. A composition according to claim 9, wherein viscosity is at least 3000 cps.

11. An extruded toilet bar composition comprising 5-50% chips comprising:

(a) 40% to about 80% by wt. of chip composition of a carrier selected from the group consisting of:

(1) hydrophobically modified polyalkylene glycol having MW of about 4000 - 25,000, wherein the compound has formula

(AG)m-R or R-(AG)m-R,

wherein AG is alkylene glycol monomer unit, m > 50 and R is attached hydrophilic group;

(2) polyoxyethylene-polyoxypropylene copolymer having MW about 4000 to 25,000;

- 43 -

(3) mixtures of (1) and (2); and

(4) mixture of (1) and/or (2) with polyalkylene glycol having MW greater than 4000 to 20,000.

(b) 10% to 40% by wt. chip composition benefit agent;

5 (c) 0.01 to 10% by wt. chip composition thickener;

(d) 0 to 10% by wt. chip composition water; and

(e) 0% to 15% by wt. chip composition, i.e.,
structuring aid/filler selected from the group
consisting of C₈ to C₂₄ fatty acids or ester
10 derivatives, and C₈ to C₂₄ alcohols or ether
derivatives,

wherein the viscosity of the thickened carrier is equal
to or greater than 800 cps,

wherein the viscosity of the thickened carrier is equal
15 to or greater than 800 cps, and 80-60% chips comprising 5 to
95% of a surfactant system wherein the surfactant is selected
from the group consisting of soap, anionic surfactant,
nonionic surfactant, amphoteric surfactant, cationic
surfactant and mixtures thereof.

20

12. A toilet bar composition according to claim 11,
wherein the surfactant system comprises

(a) a first anionic surfactant; and

(b) a second surfactant selected from the group
25 consisting of a second anionic different from the
first, a nonionic, an amphoteric and mixtures
thereof.

13. A toilet bar composition according to claim 12,
30 wherein the first anionic is acyl isethionate.

- 44 -

14. A toilet bar composition according to claim 13, wherein the isethionate forms 10% to 70% of the final bar composition.

5 15. A toilet bar composition according to any of claims 12-14, wherein the second surfactant is sulfosuccinate.

16. A toilet bar composition according to any of claims 12-15, wherein the second surfactant is betaine.

10

17. A toilet bar composition according to claim 16, wherein the betaine is amidococoylbetaine.

15 18. A toilet bar composition according to any of claims 12-17, wherein the second surfactant comprises a mixture of sulfosuccinate and betaine.

19. A toilet bar composition according to any of claims 12-18, wherein the alkylene glycol carrier is propylene glycol.

20

20. A toilet bar composition according to any of claims 12-19, wherein the benefit agent comprises silicone, petrolatum, maleated soybean oil, or sunflower seed oil, or mixtures thereof.

25

21. A toilet bar composition according to any of claims 12-20, wherein thickener comprises a fumed silica or a water soluble starch.

30

22. A process of forming a chip comprising:

- (a) 40% to about 80% by wt. of chip composition of a carrier selected from the group consisting of:

- 45 -

- (1) hydrophobically modified polyalkylene glycol having MW of about 4000 - 25,000, wherein the compound has formula



wherein AG is alkylene glycol monomer unit, $m > 50$ and R is attached hydrophilic group;

- (2) polyoxyethylene-polyoxypropylene copolymer having MW about 4000 to 25,000;

- (3) mixtures of (1) and (2); and

- (4) mixture of (1) and/or (2) with polyalkylene glycol having MW greater than 4000 to 20,000.

(b) 10% to 40% by wt. chip composition benefit agent;

(c) 0.01 to 10% by wt. chip composition thickener;

(d) 0 to 10% by wt. chip composition water; and

(e) 0% to 15% by wt. chip composition, i.e., structuring aid/filler selected from the group consisting of C_8 to C_{24} fatty acids or ester derivatives, and C_8 to C_{24} alcohols or ether derivatives,

wherein the viscosity of the thickened carrier is equal to or greater than 800 cps.

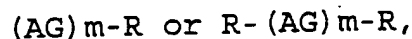
23. A process according to claim 22, wherein thickener comprises fumed silica or water soluble starch.

24. A method of enhancing deposition of benefit agent from a bar composition without compromising processing which method comprises using in said bar compositions about 20% to 40% chips comprising:

- (a) 40% to about 80% by wt. of chip composition of a carrier selected from the group consisting of:

- 46 -

- (1) hydrophobically modified polyalkylene glycol having MW of about 4000 - 25,000, wherein the compound has formula



wherein AG is alkylene glycol monomer unit,
 $m > 50$ and R is attached hydrophilic group;

- (2) polyoxyethylene-polyoxypropylene copolymer having MW about 4000 to 25,000;

- (3) mixtures of (1) and (2); and

- (4) mixture of (1) and/or (2) with polyalkylene glycol having MW greater than 4000 to 20,000.

(b) 10% to 40% by wt. chip composition benefit agent;

(c) 0.01 to 10% by wt. chip composition thickener;

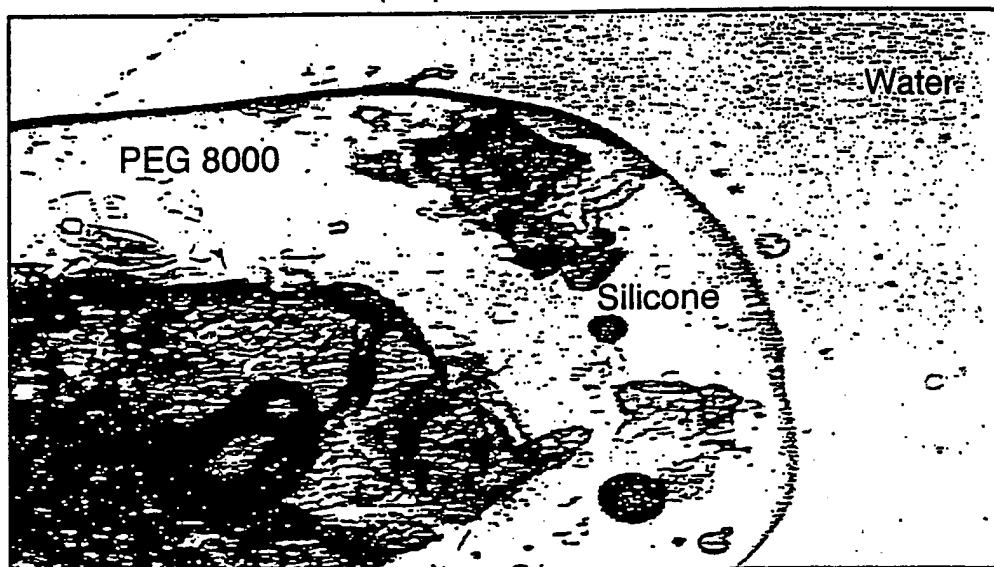
(d) 0 to 10% by wt. chip composition water; and

(e) 0% to 15% by wt. chip composition, i.e., structuring aid/filler selected from the group consisting of C_8 to C_{24} fatty acids or ester derivatives, and C_8 to C_{24} alcohols or ether derivatives,

wherein the viscosity of the thickened carrier is equal to or greater than 800 cps.

Fig.1.

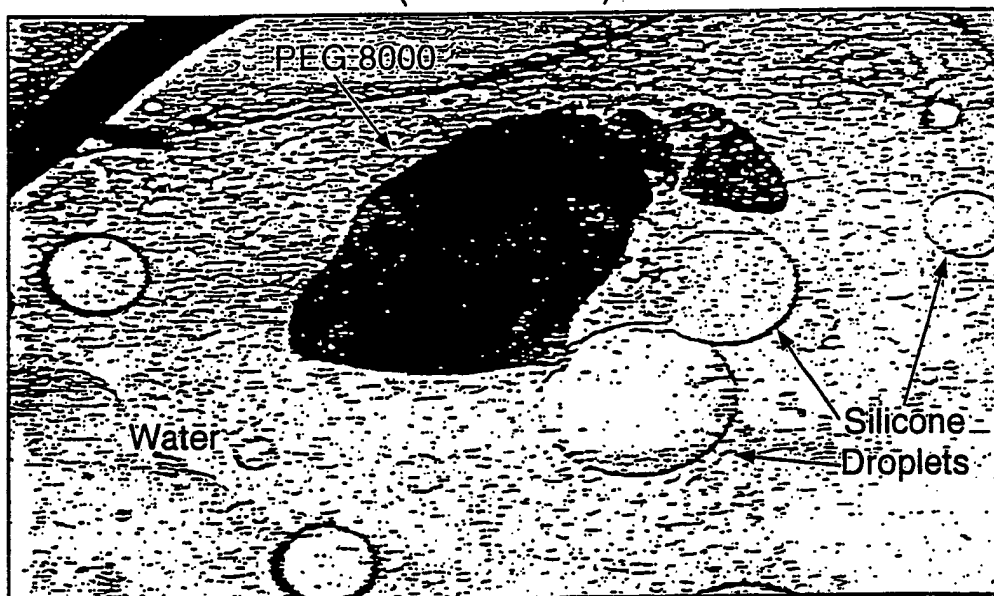
Chip of Visscher et al.
(Unprocessable)



■ Silicon coats PEG chip

Fig.2.

Chip of Invention
(Processable)



- PEG traps PDMS as droplets upon crystallisation
- PEG chip goes into solution when flooded
- Individual PDMS droplets released

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 98/06989

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C11D17/00 C11D3/37 C11D17/06 C11D3/18 C11D3/12
 C11D3/22 C11D11/00 C11D10/04 C11D1/94 //C11D1:04,
 C11D1:12,C11D1:90

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C11D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,A	WO 97 49381 A (UNILEVER) 31 December 1997 31 December 1997 see claims see example 1 see page 7, line 13 - page 8, line 28	1-24
P,A	WO 98 44086 A (UNILEVER) 8 October 1998 cited in the application 8 October 1998 see claims see examples 2,3 see page 7, line 23 - page 9, line 5 see page 23, line 6 - line 9 --- -/--	1-21



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

3 March 1999

Date of mailing of the international search report

22/03/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

Neys, P

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 98/06989

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4 927 557 A (REVIS J.W. ET AL.) 22 May 1990 see claims 1-3,5-7,9 see example 1 see column 1, line 54 - line 64 see column 2, line 22 - line 36 see column 4, line 1 - line 16 ----	1-3,7
A	WO 97 40132 A (UNILEVER) 30 October 1997 30 October 1997 see claims 1-8 see examples ----	1,2,11, 19,22,24
A	US 3 766 097 A (ROSMARIN P.) 16 October 1973 see claims 1-8 ----	1,2,11, 22,24
A	US 3 312 627 A (HOOKER D.T.) 4 April 1967 see claims 1,2 see examples ----	1,2,11, 22,24
A	WO 94 17172 A (UNILEVER) 4 August 1994 4 August 1994 see claims see examples see page 19, line 8 - line 19 ----	1-6, 11-18,20
A	WO 92 08444 A (PROCTER & GAMBLE) 29 May 1992 cited in the application see claims see examples see page 17, line 7 - line 12 -----	1,11-18, 20

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 98/06989

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 9749381	A	31-12-1997	US	5783536 A	21-07-1998
			AU	2093197 A	14-01-1998
WO 9844086	A	08-10-1998	AU	7640898 A	22-10-1998
US 4927557	A	22-05-1990	CA	1297251 A	17-03-1992
			GB	2204825 A	23-11-1988
			JP	1047436 A	21-02-1989
			JP	2672326 B	05-11-1997
WO 9740132	A	30-10-1997	US	5795852 A	18-08-1998
			AU	2025097 A	12-11-1997
			CA	2248079 A	30-10-1997
US 3766097	A	16-10-1973	NONE		
US 3312627	A	04-04-1967	NONE		
WO 9417172	A	04-08-1994	AU	5884594 A	15-08-1994
			DE	69413320 D	22-10-1998
			DE	69413320 T	25-02-1999
			EP	0631615 A	04-01-1995
			ES	2123760 T	16-01-1999
			US	5661120 A	26-08-1997
WO 9208444	A	29-05-1992	US	5154849 A	13-10-1992
			AT	141155 T	15-08-1996
			AU	9085291 A	11-06-1992
			CA	2095023 A	17-05-1992
			CN	1062373 A, B	01-07-1992
			DE	69121416 D	19-09-1996
			DE	69121416 T	20-02-1997
			DK	557423 T	09-12-1996
			EG	19614 A	30-08-1995
			EP	0557423 A	01-09-1993
			ES	2090592 T	16-10-1996
			FI	932200 A	14-05-1993
			GR	3020816 T	30-11-1996
			JP	6502646 T	24-03-1994
			MX	9102073 A, B	01-06-1992
			NZ	240610 A	27-04-1994
			PT	99531 A	30-10-1992

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ BLACK BORDERS
- ☐ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES
- ☐ FADED TEXT OR DRAWING
- ☒ BLURRED OR ILLEGIBLE TEXT OR DRAWING
- ☐ SKEWED/SLANTED IMAGES
- ☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS
- ☐ GRAY SCALE DOCUMENTS
- ☒ LINES OR MARKS ON ORIGINAL DOCUMENT
- ☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY
- ☐ OTHER: _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.